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STRUCTURE FILE UPDATES: 21 OCT 2003 HIGHEST RN 607679-40-3 DICTIONARY FILE UPDATES: 21 OCT 2003 HIGHEST RN 607679-40-3

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2003

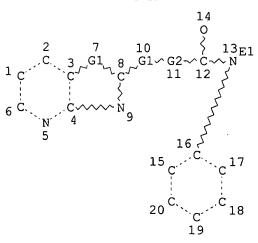
Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

*** YOU HAVE NEW MAIL ***
'.REGISTRY' IS DEFAULT FORMAT FOR 'REGISTRY' FILE

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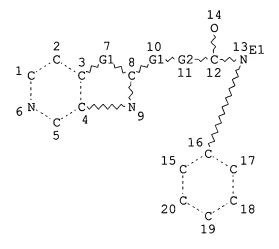


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STEREO ATTRIBUTES: NONE

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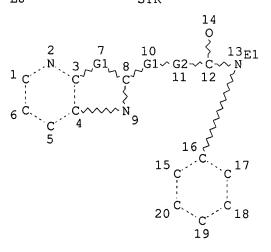
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100.0% PROCESSED 568 ITERATIONS

8 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

PROJECTED ITERATIONS: 9931 TO 12789

PROJECTED ANSWERS: 8 TO 329

L12

8 SEA SSS SAM L8 OR L9 OR L10 OR L11

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L12 8 ANSWERS REGISTRY COPYRIGHT 2003 ACS on STN
IN Benzoic acid, 4-[[((3-methyl-3H-imidazo(4,5-b)pyridin-2-yl)thio]acetyl]amino]-, methyl ester (9CI)
MF C17 H16 N4 O3 S

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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

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L13 ANSWER 53 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 603093-75-0 REGISTRY

CN Acetamide, 2-[(3-cyclohexyl-3H-imidazo[4,5-b]pyridin-2-yl)thio]-N-(2,4-dimethoxyphenyl)- (9CI) (CA INDEX NAME)

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MF C22 H26 N4 O3 S

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L13 ANSWER 50 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 603093-83-0 REGISTRY

CN Acetamide, 2-[(3-cyclohexyl-3H-imidazo[4,5-b]pyridin-2-yl)thio]-N-(2-methoxy-5-methylphenyl)- (9CI) (CA INDEX NAME)

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MF C22 H26 N4 O2 S

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=> d 37 sub bib abs

L13 ANSWER 37 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 603094-08-2 REGISTRY

CN Propanamide, N-[3-[[[[3-(phenylmethyl)-3H-imidazo[4,5-b]pyridin-2-yl]thio]acetyl]amino]phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C24 H23 N5 O2 S

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L13 ANSWER 54 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 590395-43-0 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-N-(2-iodophenyl)- (9CI) (CA INDEX NAME)

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MF C14 H11 I N4 O S

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L13 ANSWER 57 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 442660-68-6 REGISTRY

CN Acetamide, N-[4-(chlorodifluoromethoxy)phenyl]-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C15 H11 C1 F2 N4 O2 S

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RN 441782-62-3 REGISTRY

CN Acetamide, N-(2,5-dichlorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H10 C12 N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 59 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 441782-61-2 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-N-(2-methoxyphenyl)-(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C15 H14 N4 O2 S

SR Chemical Library

LC STN Files: CHEMCATS

$$\begin{array}{c|c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ &$$

L13 ANSWER 60 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 441316-85-4 REGISTRY

CN Acetamide, N-(2-chlorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H11 C1 N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

=> d 112 sub bib abs

L13 ANSWER 112 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 53052-28-1 REGISTRY

CN Acetamide, 2-(oxazolo[4,5-b]pyridin-2-ylthio)-N-phenyl- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Oxazolo[4,5-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C14 H11 N3 O2 S

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

AN 81:13482 CA

TI 2-Thio[4,5-b]oxazolopyridines

PA Ferlux-Chimie S. A.

SO Fr. Demande, 23 pp.

CODEN: FRXXBL

DT Patent

LA French

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	FR 2190426	A1	19740201	FR 1972-23119	19720627
	FR 2190426	В1	19750620		

PRAI FR 1972-23119 19720627

GI For diagram(s), see printed CA Issue.

AB Oxazolopyridinethio-acetates I (R = H, Me, Rl = OH, its amine salts, OEt; R = H, Rl = alkoxy, aryloxy, amino) (34 compds.) were prepd. by treating 2-mercaptooxazolo[4,5-b]pyridine with ClCR2COR1. I (R = H, Rl = OH.H2NCHMe2) gave 60% redn. in the writhing syndrome in mice at 300 mg/kg orally. I [R = H, Rl = OH.-HN(CHMe2)2] was hypotensive in the chloralosed dog at 20 mg/kg i.v. I (R = Me, Rl = OH) was choleretic in rats at 200 mg/kg orally.

=> d 61-65 sub bib abs

L13 ANSWER 61 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 439946-03-9 REGISTRY

CN Acetamide, N-(4-bromophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H11 Br N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

$$\begin{array}{c|c} & & & \\ &$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 62 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 432020-81-0 REGISTRY

CN Acetamide, N-(3,4-dimethylphenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C16 H16 N4 O S

SR Chemical Library

L13 ANSWER 63 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 432012-83-4 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-N-phenyl- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H12 N4 O S

SR Chemical Library

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 64 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 432003-15-1 REGISTRY

CN Benzoic acid, 4-[[(1H-imidazo[4,5-b]pyridin-2-ylthio)acetyl]amino]-, 1-methylethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C18 H18 N4 O3 S

SR Chemical Library

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 65 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 431909-18-1 REGISTRY

CN Acetamide, N-(5-chloro-2-methoxyphenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C15 H13 C1 N4 O2 S

SR Chemical Library

$$\begin{array}{c|c} & & & C1 \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & \\ & & \\ & & \\ & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ & & \\ &$$

=> d 66-70 sub bib abs

L13 ANSWER 66 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 429648-25-9 REGISTRY

CN Acetamide, N-(4-chloro-3-fluorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H10 C1 F N4 O S

SR Chemical Library

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PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 67 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 428859-87-4 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-N-[4-(1-methylpropyl)phenyl]- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C18 H20 N4 O S

SR Chemical Library

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 68 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 401611-89-0 REGISTRY

CN Acetamide, N-(3-fluorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H11 F N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 69 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 400065-85-2 REGISTRY

CN Acetamide, N-1,3-benzodioxol-5-yl-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)-(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C15 H12 N4 O3 S

SR Chemical Library

LC STN Files: CHEMCATS

$$\begin{array}{c|c} N & N & O & O \\ \hline N & N & S - CH_2 - C - NH & O \\ \hline \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 70 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN , 353761-68-9 REGISTRY

CN Acetamide, N-(2,4-difluorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H10 F2 N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

$$\begin{array}{c|c} & & & \\ &$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

\Rightarrow d 71-75 sub bib abs

L13 ANSWER 71 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 335208-80-5 REGISTRY

CN Benzoic acid, 4-[[(1H-imidazo[4,5-b]pyridin-2-ylthio)acetyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C17 H16 N4 O3 S

SR Chemical Library

LC STN Files: CHEMCATS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 72 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 335208-76-9 REGISTRY

CN Acetamide, N-(4-fluorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H11 F N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 73 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 335207-35-7 REGISTRY

CN Acetamide, N-(2,4-dimethoxyphenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)(9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C16 H16 N4 O3 S

SR Chemical Library

LC STN Files: CHEMCATS

L13 ANSWER 74 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 333415-44-4 REGISTRY

CN Acetamide, N-(2-fluorophenyl)-2-(1H-imidazo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C14 H11 F N4 O S

SR Chemical Library

LC STN Files: CHEMCATS

$$\begin{array}{c|c} N & N & S-CH_2-C-NH \\ \hline \\ NH & \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L13 ANSWER 75 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 213685-58-6 REGISTRY

CN Hexanamide, N-[2,6-bis(1-methylethyl)phenyl]-6-(oxazolo[4,5-b]pyridin-2-ylamino)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C24 H32 N4 O2

SR CA

LC STN Files: CA, CAPLUS, TOXCENTER, USPATFULL

$$\begin{array}{c|c} O & i-Pr \\ \hline O & N \\ \hline N & N \\ \hline \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

AN 129:275910 CA

TI Preparation of novel anilide compounds as acyl CoA cholesterol acyltransferase inhibitors

```
Shibuya, Kimiyuki; Kawamine, Katsumi; Sato, Yukihiro; Edano, Toshiyuki;
IN
     Edano, Toshiyuki; Hirata, Mitsuteru; Ozaki, Chiyoka
PΑ
     Kowa Company, Ltd., Japan
SO
     PCT Int. Appl., 130 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     Japanese
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                         APPLICATION NO.
                                                           DATE
                    ____
                          -----
                                          -----
                                                           _____
                                         WO 1998-JP1337
    WO 9842680
                     A1 19981001
                                                           19980325
PΙ
        W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
            DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
            KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO,
            NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA,
            UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
             FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM,
             GA, GN, ML, MR, NE, SN, TD, TG
     AU 9865176
                           19981020
                                          AU 1998-65176
                                                           19980325
                     A1
     EP 1020451
                      A1
                           20000719
                                          EP 1998-911008
                                                           19980325
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
        R:
             IE, FI
     US 6362208
                      В1
                           20020326
                                          US 1999-381850
                                                           19991206
     US 2002099074
                                          US 2001-34669
                      Α1
                           20020725
                                                           20011219
     US 2003087928
                      A1
                           20030508
                                          US 2002-79641
                                                           20020220
PRAI JP 1997-90146
                      19970325
     WO 1998-JP1337
                      19980325
     US 1999-381850
                     19991206
GΙ
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$$\begin{array}{c|c}
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$$\begin{array}{c|c}
 & Y \\
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$$\begin{array}{c|c}
 & Y \\
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\end{array}$$

The title compds. [I; A represents a divalent residue of substituted AB benzene, benzene fused with an optionally substituted heterocycle, pyridine, cyclohexane or naphthalene, or CH:CH; Ar represents optionally substituted aryl; X represents NH, oxygen or sulfur; Y represents NR4, oxygen, sulfur, sulfoxy or sulfone; Z represents a single bond or NR5; R4, R5 represent each hydrogen, lower alkyl, (un)substituted aryl, silylated lower alkyl; n is an integer of from 0 to 15] are prepd. I, possessing acyl CoA cholesterol acyltransferase (ACAT) inhibitory activity, are useful for prevention and treatment of arteriosclerosis, hyperlipemia, angina pectoris, artery tumor, ischemic heart and intestine diseases. Thus, 2-mercaptooxazolon[4,5-b]pyridine was reacted with 6-bromo-N-(2,6-diisopropylphenyl)hexeneamide in the presence of K2CO3 and 18-crown-6 to give 27% the title compd. (II), which showed IC50 of 0.14 .mu.M against ACAT from small intestine when tested with rabbit. RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 76 96 sub bib abs

ANSWER 76 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 213684-77-6 REGISTRY

CN Hexanamide, N-[2,6-bis(1-methylethyl)phenyl]-6-(thiazolo[5,4-b]pyridin-2ylthio) - (9CI) (CA INDEX NAME)

FS 3D CONCORD

C24 H31 N3 O S2 MF

CA SR

STN Files: CA, CAPLUS, TOXCENTER, USPATFULL LC

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

ΑN 129:275910 CA

ΤI Preparation of novel anilide compounds as acyl CoA cholesterol acyltransferase inhibitors

IN Shibuya, Kimiyuki; Kawamine, Katsumi; Sato, Yukihiro; Edano, Toshiyuki; Edano, Toshiyuki; Hirata, Mitsuteru; Ozaki, Chiyoka

PΑ Kowa Company, Ltd., Japan

SO PCT Int. Appl., 130 pp.

CODEN: PIXXD2

DT Patent

T.A Jananaca

LA FAN.	-	panes 1	е															
			NO.		KI	ND	DATE		APPLICATION NO. DATE									
ΡI	WO	9842	680		Α	1	1998	1001		M	0 19	98-J	P133	7	1998	0325		
		W:	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,	DE,
			DK,	EE,	ES,	FI,	GB,	GE,	GH,	GM,	GW,	ΗU,	ID,	IL,	IS,	JP,	ΚE,	KG,
			KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO,
			NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	UA,
			UG,	US,	UZ,	VN,	YU,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM	
		RW:	GH,	GM,	KE,	LS,	MW,	SD,	SZ,	UG,	ZW,	ΑT,	BE,	CH,	DE,	DK,	ES,	FI,
			FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ΒJ,	CF,	CG,	CI,	CM,
			GA,	GN,	ML,	MR,	NE,	SN,	TD,	TG								
	ΑU	9865	176		Α	1	1998	1020		A	U 19:	98-6	5176		1998	0325		
	EΡ	1020	451		Α	1	2000	0719		E	P 19	98-9	1100	8	1998	0325		
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙT,	LI,	LU,	NL,	SE,	MC,	PT,
			ΙE,	FI														
	US	6362	208		В	1	2002	0326		U	S 19	99-3	31850	0	1999	1206		
	US	2002	0990	74	Α	1	2002	0725		U	S 20	01-3	4669		2001	1219		
	US	2003	0879	28	Α	1	2003	0508		U.	S 20	02-7	9641		2002	0220		
PRAI	JP	1997	-901	46	19	9703	25											
	WO	1998	-JP1	337	19	9803	25											
	US	1999	-381	850	19	9912	06											

$$\begin{array}{c|c}
O & Y \left[CH_2 \right] z - CO - NH - Ar \\
\hline
S & I
\end{array}$$

$$\begin{array}{c|c}
 & i-Pr \\
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 & \downarrow$$

The title compds. [I; A represents a divalent residue of substituted benzene, benzene fused with an optionally substituted heterocycle, pyridine, cyclohexane or naphthalene, or CH:CH; Ar represents optionally substituted aryl; X represents NH, oxygen or sulfur; Y represents NR4, oxygen, sulfur, sulfoxy or sulfone; Z represents a single bond or NR5; R4, R5 represent each hydrogen, lower alkyl, (un)substituted aryl, silylated lower alkyl; n is an integer of from 0 to 15] are prepd. I, possessing acyl CoA cholesterol acyltransferase (ACAT) inhibitory activity, are useful for prevention and treatment of arteriosclerosis, hyperlipemia, angina pectoris, artery tumor, ischemic heart and intestine diseases. Thus, 2-mercaptooxazolon[4,5-b]pyridine was reacted with 6-bromo-N-(2,6-diisopropylphenyl)hexeneamide in the presence of K2CO3 and 18-crown-6 to give 27% the title compd. (II), which showed IC50 of 0.14 .mu.M against ACAT from small intestine when tested with rabbit.

RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 96 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 202123-74-8 REGISTRY

CN Acetamide, N-[2'-(aminosulfonyl)[1,1'-biphenyl]-4-yl]-2-(1H-imidazo[4,5-c]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C20 H17 N5 O3 S2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)

2 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

AN 130:252241 CA

TI Preparation of amidinoindoles and analogs as factor Xa inhibitors

IN Dominguez, Celia; Han, Qi; Duffy, Daniel Emmett; Park, Jeongsook Maria; Quan, Mimi Lifen; Rossi, Karen Anita; Wexler, Ruth Richmond

PA Dupont Pharmaceuticals Company, USA

SO U.S., 46 pp. CODEN: USXXAM

DT Patent

LA English

FAN.CNT 1

111111	PATENT NO.	KIND	DATE	API	PLICATION NO.	DATE		
ΡI	US 5886191	A	19990323	US	1997-916736	19970818		
	US 6043257	Α	20000328	US	1998-176037	19981021		
PRAI	US 1997-916736	19970	818					
GT								

$$\mathbb{R}^{2}$$
 \mathbb{R}^{2}
 \mathbb{R}^{2}
 \mathbb{R}^{1}

Title compds., e.g., I [R1 = H or Me; R2 = (CH2)nZZ1R; R = C(:NH)NH2, CH2Ph, C6H4(SO2NHR4)-2, etc.; R3 = C(:NH)NH2, cyano, etc.; R4 = alkyl; Z = CO, CONH, etc.; Z1 = C6H4, CH2C6H4, pyridine-2,4-diyl, etc.; n = 0 or 1; dashed line = optional addnl. bond] were prepd. as factor Xa inhibitors (no data). Thus, 5-cyanoindole was acylated by (COCl)2 and the product converted in 3 steps to 5-cyanoindole-3-acetic acid which was amidated by 4-(2-aminosulfonylphenyl)-2-pyridinamine to give, in 2 addnl. steps, I [R1 = H, R2 = CH2CONHZ1C6H4(SO2NH2)-2, R3 = C(:NH)NH2, Z1 = pyridine-2,4-diyl, dashed line = bond].

RE.CNT 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

REFERENCE 2

AN 128:128015 CA

TI Preparation of amidinoindoles and amidinoazoles as inhibitors of factor Xa and of thrombin

IN Dominguez, Celia; Han, Qi; Duffy, Daniel Emmett; Park, Jeongsook Maria; Quan, Mimi Lifen; Rossi, Karen Anita; Wexler, Ruth Richmond

PA Du Pont Merck Pharmaceutical Co., USA

SO PCT Int. Appl., 176 pp. CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PA:	CENT	NO.		KI	KIND DATE					APPLICATION NO.				DATE				
										_									
PI	WO 9801428				A	1	19980115			WO 1997-US11325					19970630				
		W:	AM,	ΑU,	AZ,	BR,	BY,	CA,	CN,	CZ,	EE,	HU,	IL,	JP,	KG,	KR,	ΚZ,	LT,	
			LV,	MD,	MX,	NO,	NZ,	PL,	RO,	RU,	SG,	SI,	SK,	ТJ,	TM,	UA,	VN,	AM,	
			ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM									
		RW:	ΑT,	BE,	CH,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	ΙT,	LU,	MC,	NL,	PT,	SE
	ÇA	2259	573		\mathbf{A}	A	1998	0115		C	A 19	97-2	2595	73	1997	0630			
	ΑU	9736	456		A	1	1998	0202		Αl	U 19	97-3	6456		1997	0630			
	EΡ	9601	02		A	1	1999	1201		E.	P 19	97-9	3321	4	1997	0630	•		
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	PT,	ΙE	
	NZ	3336	96		Α		2000	0623		N.	Z 19	97 - 3	3369	6	1997	0630			

PRAI US 1996-676766 19960708 US 1997-49519P 19970613 WO 1997-US11325 19970630

GI

AB The title compds. [I; W, W3 = CH, N; W1, W2 = C, CH, N (provided that one of W1 and W2 is C(C(=NH)NH2) and at most two of W, W1, W2, and W3 are N); one of D, Da = H, C1-4 alkoxy, CN, etc. and the other is absent; one of Ja and Jb is substituted by -(CH2)n-Z-A-B; J, Ja, Jb combine to form an arom. heterocyclic system contg. from 1-2 heteroatoms (N, O, and S), a heterocyclic ring wherein Jb = N and J and Ja = (un)substituted CH2, a heterocyclic ring wherein Jb = CH, J = (un)substituted NH and Ja = (un) substituted CH; Z = CH:CH, SO2CH2, etc.; A = (un) substituted PhCH2, PhCH2CH2, etc.; B = C3-6 alkyl, (un) substituted PhCH2, 5-10 membered heterocyclic system, etc.], useful as inhibitors of factor Xa or thrombin, were prepd. and formulated. Thus, reaction of 5-cyanoindole-1-acetic acid with 4-benzylpiperidine followed by treatment of the resulting 1-(4-benzylpiperidinocarbonyl)methyl-5-cyanoindole with HCl(q) in MeOH, and then with (NH4)2CO3 in MeOH afforded the title compd. II. Some compds. I were evaluated and showed Ki of < 5 .mu.M against thrombin. RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 97 98 106 sub bib abs

L13 ANSWER 97 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 202123-73-7 REGISTRY

CN Acetamide, N-[2'-[[(1,1-dimethylethyl)amino]sulfonyl][1,1'-biphenyl]-4-yl]-2-(1H-imidazo[4,5-c]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

FS 3D CONCORD

MF C24 H25 N5 O3 S2

SR CA

LC STN Files: CA, CAPLUS, USPATFULL

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

2 REFERENCES IN FILE CA (1907 TO DATE)

REFERENCE 1

AN 130:252241 CA

TI Preparation of amidinoindoles and analogs as factor Xa inhibitors

IN Dominguez, Celia; Han, Qi; Duffy, Daniel Emmett; Park, Jeongsook Maria; Quan, Mimi Lifen; Rossi, Karen Anita; Wexler, Ruth Richmond

PA Dupont Pharmaceuticals Company, USA

SO U.S., 46 pp. CODEN: USXXAM

DT Patent

LA English

FAN CNT 1

r AN. CN	AT T					
P	PATENT NO.	KIND	DATE	AP	PLICATION NO.	DATE
_						
PI U	JS 5886191	Α	19990323	US	1997-916736	19970818
Ü	JS 6043257	Α	20000328	US	1998-176037	19981021
PRAI U	JS 1997-916736	19970	818			

$$\mathbb{R}^{3}$$
 \mathbb{N}
 \mathbb{R}^{1}
 \mathbb{I}

Title compds., e.g., I [R1 = H or Me; R2 = (CH2)nZZ1R; R = C(:NH)NH2, CH2Ph, C6H4(SO2NHR4)-2, etc.; R3 = C(:NH)NH2, cyano, etc.; R4 = alkyl; Z = CO, CONH, etc.; Z1 = C6H4, CH2C6H4, pyridine-2,4-diyl, etc.; n = 0 or 1; dashed line = optional addnl. bond) were prepd. as factor Xa inhibitors (no data). Thus, 5-cyanoindole was acylated by (COCl)2 and the product converted in 3 steps to 5-cyanoindole-3-acetic acid which was amidated by 4-(2-aminosulfonylphenyl)-2-pyridinamine to give, in 2 addnl. steps, I [R1 = H, R2 = CH2CONHZ1C6H4(SO2NH2)-2, R3 = C(:NH)NH2, Z1 = pyridine-2,4-diyl, dashed line = bond].

RE.CNT 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

REFERENCE 2

AN 128:128015 CA

TI Preparation of amidinoindoles and amidinoazoles as inhibitors of factor Xa and of thrombin

IN Dominguez, Celia; Han, Qi; Duffy, Daniel Emmett; Park, Jeongsook Maria;
Quan, Mimi Lifen; Rossi, Karen Anita; Wexler, Ruth Richmond

PA Du Pont Merck Pharmaceutical Co., USA

SO PCT Int. Appl., 176 pp. CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.			KI	ND	DATE			Α	PPLI	CATI	ON N	ο.	DATE					
										_									
ΡI	WO	9801	428		Α	1	1998	0115		W	0 19	97 - U	S113	25	1997	0630			
		W:	AM,	ΑU,	ΑZ,	BR,	BY,	CA,	CN,	CZ,	EE,	ΗU,	IL,	JP,	KG,	KR,	ΚŹ,	LT,	
			LV,	MD,	MX,	NO,	NZ,	PL,	RO,	RU,	SG,	SI,	SK,	TJ,	TM,	UA,	VN,	AM,	
			ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM									
		RW:	AT,	BE,	CH,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE

The title compds. [I; W, W3 = CH, N; W1, W2 = C, CH, N (provided that one AΒ of W1 and W2 is C(C(=NH)NH2) and at most two of W, W1, W2, and W3 are N); one of D, Da = H, C1-4 alkoxy, CN, etc. and the other is absent; one of Ja and Jb is substituted by -(CH2)n-Z-A-B; J, Ja, Jb combine to form an arom. heterocyclic system contg. from 1-2 heteroatoms (N, O, and S), a heterocyclic ring wherein Jb = N and J and Ja = (un)substituted CH2, a heterocyclic ring wherein Jb = CH, J = (un)substituted NH and Ja = (un) substituted CH; Z = CH:CH, SO2CH2, etc.; A = (un) substituted PhCH2, PhCH2CH2, etc.; B = C3-6 alkyl, (un) substituted PhCH2, 5-10 membered heterocyclic system, etc.], useful as inhibitors of factor Xa or thrombin, were prepd. and formulated. Thus, reaction of 5-cyanoindole-1-acetic acid with 4-benzylpiperidine followed by treatment of the resulting 1-(4-benzylpiperidinocarbonyl)methyl-5-cyanoindole with HCl(q) in MeOH, and then with (NH4)2CO3 in MeOH afforded the title compd. II. Some compds. I were evaluated and showed Ki of < 5 .mu.M against thrombin.

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 98 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-70-6 REGISTRY

CN Acetamide, 2-(thiazolo[5,4-b]pyridin-2-ylthio)-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H10 F3 N3 O S2

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

AN 99:158411 CA

TI Thiazolopyridinylthioalkanamides

PA Otsuka Pharmaceutical Factory, Inc., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

CAN.	CNII						
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
ΡI	JP 58116489	A2	19830711	JP 1981-215860	19811228		
	JP 62037038	B4	19870810				
PRAI	JP 1981-215860	19811	.228				

$$R \xrightarrow{N} SZCONR^{1}R^{2}$$

$$I$$

$$N$$

$$S$$

$$II$$

AB The title compds. I [R = H, halo; R1,R2 = H, alkyl, cycloalkyl, alkenyl, (un)substituted Ph; Z = (un)substituted alkylene], useful as antiinflammatories, antihypertensives, analgesics, and antipyretics (no data), were prepd. Thus, refluxing 1.8 g PhCHClCONHMe with 1.6 g thiazolopyridine II, 1.5 g NaI, and 1.0 g Na2CO3 in acetone gave 1.3 g I (R = R1 = H, R2 = Me, Z = PhCH).

L13 ANSWER 106 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 75426-89-0 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-c]pyridin-2-ylthio)-N-(2-methylphenyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Imidazo[4,5-c]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H14 N4 O S

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT (*File contains numerically searchable property data)

$$S-CH_2-C-NH$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

AN 94:65548 CA

TI Synthesis and tuberculostatic activity of some derivatives of 2-mercaptoimidazo[4,5-c]pyridine

- AU Czarnocka-Janowicz, Anna; Sawlewicz, Jozef; Jakubowski, Joachim; Janowiec, Mieczysław
- CS Inst. Technol. Anal. Pharm. Prod., Sch. Med., Gdansk, 80-416, Pol.

SO Acta Poloniae Pharmaceutica (1979), 36(5), 529-37

CODEN: APPHAX; ISSN: 0001-6837

DT Journal

LA Polish

GΙ

The title compds. I, (R = OH, OMe, NH2, NHNH2, NHPh, cyclohexylamino, NPh2, NHCH2Ph, NHCH2CHMe2, N(CH2CHMe2)2, NHPr, NHC6H4Me-2, -3, and -4) were prepd. in 31-70% yields by condensing 2-mercaptoimidazo[4,5-c]pyridine (II) with ClCH2COR. An analogous condensation of II with ClCH2CN yielded 45% III, subsequently converted into the amidoxime and into aldehyde derivs. IV, (R1 = Ph, 4-ClC6H4, 2-O2NC6H4). II heated in EtOH with CH2:CHCN in presence of Et3N yielded 87% V (R = CN), which by routine methods was converted into V [R = CO2H, CO2Et, CONH2, C(:NOH)NH2, C(:NNH2)NH2]. In in vitro tests, I (R = NHPr) inhibited the growth of Mycobacterium tuberculosis H37Rv strain at 1.9 .mu.g/cm.

=> d 99-105 ide

L13 ANSWER 99 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-69-3 REGISTRY

CN Acetamide, N-(4-methylphenyl)-2-(thiazolo[5,4-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H13 N3 O S2

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 100 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-68-2 REGISTRY

CN Acetamide, 2-[(6-bromothiazolo[5,4-b]pyridin-2-yl)thio]-N-[4-(1-oxopropyl)phenyl]- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C17 H14 Br N3 O2 S2

LC STN Files: CA, CAPLUS

Br
$$S-CH_2-C-NH$$
 $C-Et$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 101 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-67-1 REGISTRY

CN Acetamide, 2-(thiazolo[5,4-b]pyridin-2-ylthio)-N-(3,4,5-trimethoxyphenyl)-(9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C17 H17 N3 O4 S2

LC STN Files: CA, CAPLUS

$$\begin{array}{c|c} & & & \\ &$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 102 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-66-0 REGISTRY

CN Butanamide, N-(2,6-dichlorophenyl)-4-(thiazolo[5,4-b]pyridin-2-ylthio)-(9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, butanamide deriv.

FS 3D CONCORD

MF C16 H13 C12 N3 O S2

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 103 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-65-9 REGISTRY

CN Butanamide, N-(2-fluorophenyl)-4-(thiazolo[5,4-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, butanamide deriv.

FS 3D CONCORD

MF C16 H14 F N3 O S2

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 104 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-64-8 REGISTRY

CN Propanamide, N-(4-nitrophenyl)-3-(thiazolo[5,4-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5,4-b]pyridine, propanamide deriv.

FS 3D CONCORD

MF C15 H12 N4 O3 S2

LC STN Files: CA, CAPLUS

$$S-CH_2-CH_2-C-NH$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 105 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 87341-63-7 REGISTRY

CN Propanamide, N-phenyl-3-(thiazolo[5,4-b)pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Thiazolo[5, 4-b]pyridine, propanamide deriv.

FS 3D CONCORD

MF C15 H13 N3 O S2

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 107-112 ide

L13 ANSWER 107 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 75426-88-9 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-c]pyridin-2-ylthio)-N-(3-methylphenyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Imidazo[4,5-c]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H14 N4 O S

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT (*File contains numerically searchable property data)

H O

$$\begin{array}{c|c} H & \\ N & \\ N & \\ \end{array}$$
 S- CH₂- C- NH Me

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 108 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 75426-86-7 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-c]pyridin-2-ylthio)-N-(4-methylphenyl)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Imidazo[4,5-c]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H14 N4 O S

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT

(*File contains numerically searchable property data)

$$\begin{array}{c|c} H & O \\ N & I \\ N & S - CH_2 - C - NH \end{array}$$

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 109 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 75426-82-3 REGISTRY

CN Acetamide, 2-(1H-imidazo[4,5-c]pyridin-2-ylthio)-N-phenyl- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN 1H-Imidazo[4,5-c]pyridine, acetamide deriv.

FS 3D CONCORD

MF C14 H12 N4 O S

LC STN Files: BEILSTEIN*, CA, CAPLUS, CASREACT

(*File contains numerically searchable property data)

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)
1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 110 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 53052-30-5 REGISTRY

CN Acetamide, 2-(oxazolo[4,5-b]pyridin-2-ylthio)-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Oxazolo[4,5-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H10 F3 N3 O2 S

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 111 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 53052-29-2 REGISTRY

CN Acetamide, N-(2-chlorophenyl)-2-(oxazolo[4,5-b]pyridin-2-ylthio)- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Oxazolo[4,5-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C14 H10 C1 N3 O2 S

LC STN Files: CA, CAPLUS

$$\begin{array}{c|c} N & N & O \\ \hline & N & S - CH_2 - C - NH \\ \hline & C1 & \end{array}$$

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

L13 ANSWER 112 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 53052-28-1 REGISTRY

CN Acetamide, 2-(oxazolo[4,5-b]pyridin-2-ylthio)-N-phenyl- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Oxazolo[4,5-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C14 H11 N3 O2 S

LC STN Files: CA, CAPLUS

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

1 REFERENCES IN FILE CA (1907 TO DATE)

1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> d 110 sub bib abs

L13 ANSWER 110 OF 112 REGISTRY COPYRIGHT 2003 ACS on STN

RN 53052-30-5 REGISTRY

CN Acetamide, 2-(oxazolo[4,5-b]pyridin-2-ylthio)-N-[3-(trifluoromethyl)phenyl]- (9CI) (CA INDEX NAME)

OTHER CA INDEX NAMES:

CN Oxazolo[4,5-b]pyridine, acetamide deriv.

FS 3D CONCORD

MF C15 H10 F3 N3 O2 S

LC STN Files: CA, CAPLUS

1 REFERENCES IN FILE CA (1907 TO DATE) 1 REFERENCES IN FILE CAPLUS (1907 TO DATE)

REFERENCE 1

81:13482 CA ΑN

ΤI 2-Thio[4,5-b]oxazolopyridines

PA Ferlux-Chimie S. A.

SO Fr. Demande, 23 pp.

CODEN: FRXXBL

DT Patent

LA French

FAN.CNT 1

	J				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	FR 2190426	A1	19740201	FR 1972-23119	19720627
	FR 2190426	В1	19750620		

PRAI FR 1972-23119 19720627

For diagram(s), see printed CA Issue.

AB Oxazolopyridinethio-acetates I (R = H, Me, R1 = OH, its amine salts, OEt; R = H, R1 = alkoxy, aryloxy, amino) (34 compds.) were prepd. by treating 2-mercaptooxazolo[4,5-b]pyridine with ClCR2COR1. I (R = H, R1 =OH.H2NCHMe2) gave 60% redn. in the writhing syndrome in mice at 300 mg/kg orally. I [R = H, R1 = OH.-HN(CHMe2)2] was hypotensive in the chloralosed dog at 20 mg/kg i.v. I (R = Me, R1 = OH) was choleretic in rats at 200 mg/kg orally.

=> save 110

ENTER NAME OR (END):s10034669/a

L10 IS NOT AN ANSWER SET

A name ending in /A can only be used to save the L-number answer set resulting from a search.

=> save 113

ENTER NAME OR (END):s10034669/a

ANSWER SET L13 HAS BEEN SAVED AS 'S10034669/A'

=> loa h

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CA SUBSCRIBER PRICE	ENTRY -4.96	SESSION -6.91

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